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Nematode-Induced Changes of Transporter Gene Expression in *Arabidopsis* Roots

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Root-knot plant-parasitic nematodes (*Meloidogyne* spp.) account for much of the damage inflicted to plants by nematodes. The feeding sites of these nematodes consist of “giant” cells, which have characteristics of transfer cells found in other parts of plants. Increased transport activity across the plasma membrane is a hallmark of transfer cells, and giant cells provide nutrition for nematodes; therefore, we initiated a study to identify the transport processes that contribute to the development and function of nematode-induced feeding sites. The study was conducted over a 4-week period, during which time the large changes in the development of giant cells were documented. The *Arabidopsis* ATH1 GeneChip was used to identify the many transporter genes that were regulated by nematode infestation. Expression of 50 transporter genes from 18 different gene families was significantly changed upon nematode infestation. Sixteen transporter genes were studied in more detail using real-time reverse-transcriptase polymerase chain reaction to determine transcript abundance in nematode-induced galls that contain giant cells and uninfested regions of the root. Certain genes were expressed primarily in galls whereas others were expressed primarily in the uninfested regions of the root, and a third group was expressed evenly throughout the root. Multiple transport processes are regulated and these may play important roles in nematode feeding-site establishment and maintenance.

Additional keywords: membrane transport, microarray.

Plant-parasitic nematodes are among the most destructive plant pathogens. Root-knot nematodes (*Meloidogyne* spp.) are capable of reproducing on over 2,000 species of plants and are responsible for approximately half of the overall nematode damage in crops (Sasser 1980). The interaction between the root and the nematode begins when an infective second-stage juvenile (J2) locates and penetrates a root behind the root cap in the zone of elongation or at rupture sites of lateral roots (Krusberg and Nielsen 1958). Upon entering the root, the J2 migrates towards the stele, where it establishes a feeding site. It is hypothesized that substances initially produced in the dor-

sal and subventral pharyngeal glands of the nematode are injected into the plant cells that surround the head of the nematode to initiate the feeding site (Hussey 1989).

Symptoms of plant infestation by root-knot nematodes include knot- or gall-like formations on the roots. These knots, or galls, inhibit the ability of the root to take up nutrients and water (Dorhout et al. 1991). Knot formation is due in large part to the development of specialized feeding cells called “giant cells”. Normally, four to eight root cells undergo a morphogenesis into a feeding structure consisting of several giant, multinucleated cells that are similar to transfer cells in that they have a cell wall labyrinth with extensions into the cytoplasm (Bird 1961; Jones and Northcote 1972; Offler et al. 2003). Characteristics of giant cells also include large numbers of mitochondria and plastids, numerous small vacuoles, and dense cytoplasm (Bird 1961; Hussey and Mims 1991; Jones and Dropkin 1976). Giant cells are metabolically very active. High levels of total protein, amino acids, glucose, glucose 6-phosphate, and ATP are found in root-knot nematode-induced giant cells (Gommers and Dropkin 1977). The increased metabolic activity of giant cells stimulates the mobilization of photosynthates from the shoot to the giant cells, where the nematode can easily access and withdraw the transferred assimilates (Bird and Loveys 1975; McClure 1977).

One of the defining features of root-knot nematode-induced giant cells that suggest they function as transfer cells is the presence of cell wall ingrowths. In marked contrast to the increase in surface area, the frequency of plasmodesmata decreases, particularly in areas of cell wall ingrowths (Huang and Maggenti 1969; Jones 1980). Due to the lack of plasmodesmata, it has been concluded that the bulk of solutes enter giant cells across the plasma membrane via apoplastic transport. The relative symplastic isolation of nematode feeding cells has been observed in several plant-parasitic nematode systems (Böckenhoff et al. 1996; Dorhout et al. 1993; Jones 1980). Therefore, it is likely that specialized membrane transport proteins are deployed to regulate the flow of nutrients into and out of giant cells, as has been demonstrated for transfer cells. A recent study on symplastic trafficking in *Heterodera schachtii*-infested *Arabidopsis* roots showed that syncytial cells are not symplastically isolated (Hoth et al. 2005). Studies using these tools have not yet been reported using *Meloidogyne incognita*; therefore, giant cells may be confirmed to be symplastically isolated. In classical plant transfer cells, transporters for sucrose, amino acids, and H⁺-ATPases have been localized to the plasma membrane of these cells and are thought to control nutrient flow into the developing seed (Harrington et al. 1997; Tegeder et al. 2000; Wang et al. 1995). To date, only a few genes that

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*The e-Xtra logo stands for “electronic extra” and indicates the HTML abstract available on-line contains a supplemental table not included in the print edition, which provides data gathered from the analysis of the transporter genes found on the ATH1 GeneChip.

encode membrane transport proteins have been demonstrated to be upregulated (Gheysen and Fenoll 2002) upon nematode infestation. These genes include the aquaporin TobRB7 (Opperman et al. 1994), a sucrose transporter (Juergensen et al. 2003), and an H⁺-ATPase (Bird and Wilson 1994).

Evidence suggesting an important role for transport proteins in nematode feeding sites led us to conduct a global analysis of the regulation of transporter gene expression after nematode infection. We used a whole-genome Affymetrix ATH1 GeneChip to compare roots from control and from root-knot nematode (*M. incognita*)-infested plants. We focused our hypotheses testing for differential gene expression on only those features encoding membrane transport proteins. This allowed us to identify transporter genes whose expression changed due to nematode infestation. Additional studies using real-time polymerase chain reaction (PCR) methods and promoter:: β -glucuronidase (GUS) constructs allowed for the tissue localization of selected genes.

RESULTS

Characterization of knot and giant-cell development.

During feeding site formation, plant cells undergo dramatic changes in size and morphology, and these changes are accompanied by concomitant rearrangements of subcellular organelles. This phenomenon has been described for several plant species. Although the infection process of *Arabidopsis* by root-knot nematodes has been described previously (Sijmons et al. 1991), detailed studies focusing on the ultrastructure of giant cells throughout their development have not been reported for *Arabidopsis*. Altered expression of membrane transport proteins occurs in the context of these cellular changes; therefore, we undertook a detailed examination of the ultrastructural changes that occur during development of root knots and giant cells in *Arabidopsis* roots. Based on these studies, we selected three timepoints representing early (1 week postinoculation), intermediate (2 weeks postinoculation), and late (4 weeks postinoculation) stages during knot and giant-cell development.

Because the penetration of the roots by nematodes was not synchronous, we selected electron microscopic images that were representative of development at each timepoint (Fig. 1). At the 1-week stage, developing giant cells generally were characterized by the presence of multiple nuclei displaying lobed morphologies (*not shown*), and large numbers of vacuolar structures were visible (Fig. 1A; arrows). No significant cell wall thickening or plasma membrane proliferation was observed in these early-stage giant cells. Many of the intermediate-stage giant cells already were expanded significantly. This expansion was accompanied by and perhaps driven by the development of large central vacuole compartments (Fig. 1B; asterisk) and numerous smaller vacuolar structures (Fig. 1B; arrows). This large central vacuolar structure was seen consistently in sections that contained giant cells at an intermediate stage of development. In addition to changes in the vacuolar membranes, complexes of Golgi and other organelles were found in increased numbers around the periphery of the cell. Many giant cells possessed thickened cell walls; however, no cell wall invaginations were visible at this intermediate stage (Fig. 1B). By week 4, most giant cells were fully developed and clearly were much larger than surrounding cells (Fig. 1C). Strikingly, despite the increased cell size of fully developed giant cells, large central vacuoles were absent, with the majority of the cell volume filled with cytoplasm and numerous smaller vacuolar structures. At this stage in development, the cell wall invaginations were evident in the fully developed giant cells, indicating a concomitant proliferation of plasma mem-

brane (Fig. 1C). In addition, we often observed nematode feeding tubes in these cells, indicating that these cells were active nematode feeding sites. By week 4, the nematodes were sedentary and unable to move, at which stage the gall and its specialized giant cells provide the nutrients required by the nematode to complete its life cycle.

Differential expression of genes on the array.

As the giant cell developed, we monitored the expression of transporter genes using the Affymetrix ATH1 GeneChip. Of the 867 transport proteins represented in the *Arabidopsis* Membrane Protein Library (AMPL) database (Ward 2001), 805 were on the array and 635 were classified as "present" in at least one replicate. Based on annotation, we found that one of these was not a transporter and therefore was excluded from further consideration. Of the 634 membrane proteins examined, 50 were found to be differentially expressed ($P < 0.016$; false discovery rate (FDR) = 0.2) in the comparison of the nematode-infested roots versus control (Table 1). The pattern of gene expression showed that there were no transporters with significant differences that were downregulated in week 1 and then upregulated in later weeks. Twenty of the genes were downregulated in all three time periods. Of the remaining 30, 26 were upregulated for both week 1 and week 2 and 16 were upregulated for all three timepoints.

In the 50 genes that showed significantly altered expression, six members of the peptide transporter/proton-coupled oligopeptide transporter family were downregulated at all timepoints and only 2 members of this family were upregulated, indicating an overall downregulation of peptide transport capacity in roots infested with root-knot nematode (Table 1). In contrast to the peptide transporters, three members of the auxin amino acid permease (AAP) family were upregulated across the timepoints, whereas only one member of this family was downregulated at week 4.

In addition to amino acid transporters, it appears that transcript levels of several genes encoding calcium transporters also changed in the nematode-infested roots (Table 1). Two P-type ATPase genes (Bonza et al. 2000; Geisler et al. 2000), *AtACA8* ($P = 0.001$) and *AtACA4* ($P = 0.016$), and a single gene encoding a CAX exchanger, *AtCAX3* ($P = 0.00006$), were significantly different between nematode-infested roots and control roots and were upregulated at either week 1 or week 2 or at both timepoints (Table 1). Although, as yet, no clear functional role can be ascribed to the members of the multidrug/oligosaccharidyl-lipid/polysaccharide flippase (MOP) superfamily of transporters (Hvorup et al. 2003), we found that three generally were upregulated and two were downregulated (Table 1).

The giant cells act as sinks for photosynthates that are used by the nematodes as a carbon source. Because sucrose is the main nonreducing sugar transported to sink tissues, we expected to find that the expression of at least one sucrose transporter gene was induced. This was the case for *AtSUC1*, which was altered ($P = 0.008$) and upregulated at all three timepoints.

A number of inorganic ion transporters generally were downregulated upon nematode infestation, including members of the zinc iron permease (ZIP), voltage-gated ion channel (VIC), and ammonium transporter (AMT) families (Table 1) (Guerinot 2000; Mäser et al. 2001; Sohlenkamp et al. 2000).

Verification of microarray data.

To confirm that mRNA of features identified by microarray analysis were indeed differentially expressed, the degree of change of 16 genes was tested using quantitative real-time reverse-transcriptase polymerase chain reaction (RT-PCR) (iCycler; BioRAD, Hercules, CA, U.S.A.) by examining the control and infested tissue for each of the three timepoints.

The 16 genes were chosen to represent members of the AAP, major facilitator super family, plasma membrane intrinsic protein (PIP), and P-type ATPase families. The raw fold changes observed on the array ranged from 0.7 to 8. The fold increase in response to nematode infestation measured by real-time RT-PCR as a function of the fold change measured by microarray analysis is shown in Figure 2. The real-time RT-PCR analysis was in good agreement with the results derived from array analysis (Fig. 2): week 1, $r^2 = 0.62$ ($P = 0.0004$); week 2, $r^2 = 0.78$ ($P < 0.0001$); and week 4, $r^2 = 0.62$; ($P < 0.0001$). The qualitative change (up- or downregulated) measured in the microarray experiment were confirmed by quantitative PCR in all cases tested. This is consistent with several studies that used real-time PCR to verify microarray data and found good correlation between the two methods (Käb et al. 2004; Paul et al. 2004; Wang et al. 2003).

Tissue specificity of transporter gene expression.

Because the GeneChip experiments were completed with RNA extracted from whole roots infested with nematodes, real-time RT-PCR experiments were carried out to determine whether selected genes were specifically expressed in regions where knots are present or were upregulated throughout

infested roots. To address this question, galls from *Arabidopsis* plants representing an additional independent biological replicate were manually excised from the root tissue. Tissue from the week-2 timepoint was chosen because knots were readily excised from the surrounding tissue, and some genes upregulated at week 1 and week 4 also were induced at week 2. Genes for this analysis were chosen based upon either of two criteria. The first criterion was significance in the initial microarray study and relevance to transfer cell function. From this, we selected *AtSUC1*, *AtACA8*, *AtAUX4/LAX3* (*At2g21050*), *AtPIP2.6*, *AtAAP1*, *AtAUX1*, *AtMRP4*, and *AtANT1* for our detailed analysis (Fig. 3). The second criterion was previous evidence about amino acid transport processes in transfer cells (Tegeder et al. 2000) and evidence for water channel and amino acid transporter gene expression in giant cells after plant nematode infection (Opperman et al. 1994; Puthoff et al. 2003). Using this criterion, we selected members of the AAP family (*AtAAP6*, *AtAAP3*, and *AtAAP7*), the ANT family (*At2g41190* and *At1g80510*), the LHT family (*AtLHT4*), and the putative aquaporin *AtPIP2.5* (Fig. 3). We also found that, by further analysis of supplemental data that was not annotated in a previous, smaller-scale microarray experiment (8K GeneChip), *AtPIP2.5* and

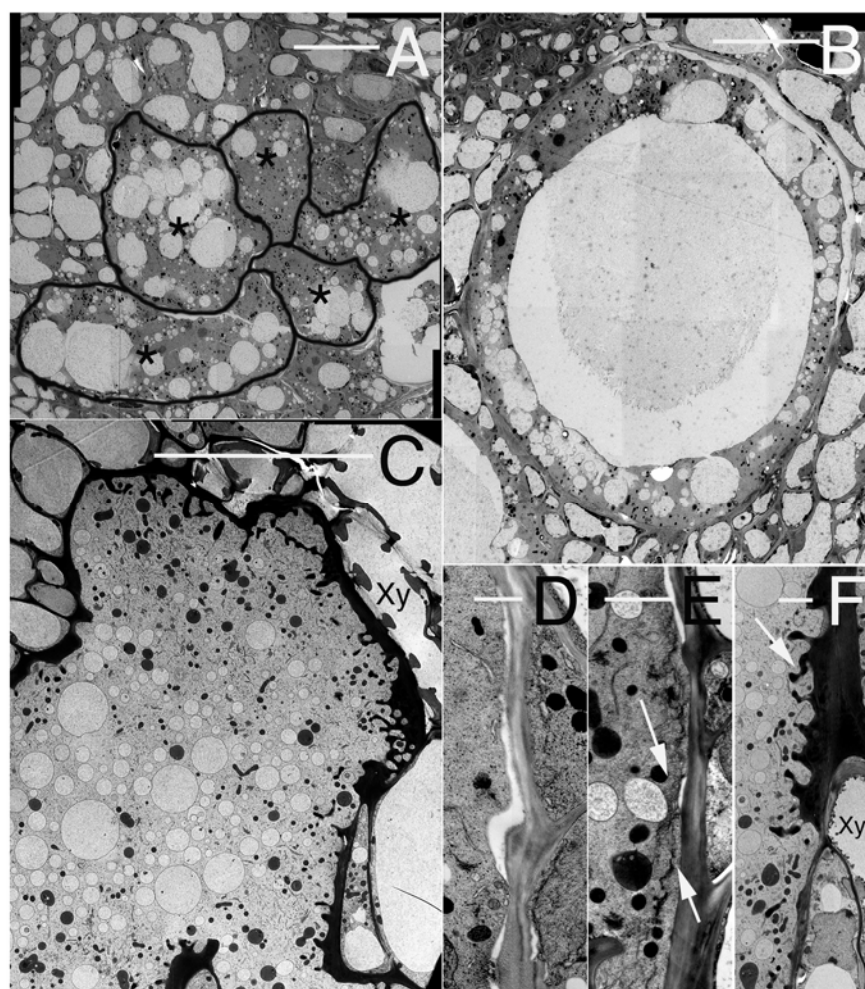


Fig. 1. Ultrastructure of freeze-substituted giant cells **A** and **D**, 1 week, **B** and **E**, 2 weeks, and **C** and **F**, 4 weeks after infestation of *Arabidopsis* roots with nematodes. Bars = 20 μ m (**A**–**C**) and 3 μ m (**D**–**F**). **A**, Formation of the feeding site was observable within 1 week after infestation. Developing giant cells (asterisks, outlined in black) contain a number of vacuoles of sizes similar to vacuoles in adjacent, nongiant cells. Many of the vacuoles were found to be in clusters with some vacuoles showing signs of fusion. **B**, Second week of infestation, giant cells have unusually large vacuoles in addition to numerous smaller vacuoles (the vacuole in this cell is 60 μ m in diameter). **C**, After 4 weeks of infestation, giant cells have lost the large vacuole and are cytoplasm rich, with a large numbers of small vacuoles (Xy, xylem element). **D** through **F**, Changes in walls of giant cells during development; in each micrograph, the giant cell is on the left side of the cell wall. At 1 week (**C**), the wall is relatively unchanged from that of nongiant cells. By the second week (**E**), a complex of ER and Golgi stacks is associated with the wall (arrows). Within 4 weeks (**F**), a transfer cell wall labyrinth develops (arrow), (Xy, xylem element).

AtAAP6 were upregulated during beet cyst nematode infestation of *Arabidopsis* roots (Puthoff et al. 2003 supplemental data).

The transcript levels of 15 selected genes were studied in galls compared with uninfected sections of root using quantitative real-time RT-PCR (Fig. 3). With the exception of *AtANT1* and *AtAAP7*, all the genes chosen for this analysis were upregulated at the week-2 timepoint (in both array and RT-PCR). These two were downregulated in both week 1 and week 2.

The detailed real-time PCR analysis of the 15 selected genes showed that they could be classified into three distinct groups. Group I included *AtPIP2.5*, *AtAAP7*, *AtSUC1*, *AtACA8*, *AtAAP6*, *AtAAP3*, and *AtAUX4/LAX3*, which were expressed at significantly higher levels ($P < 0.05$) in the galls compared

with the uninfected parts of the root (Fig. 3). In group I, the expression of aquaporin *AtPIP2.5* was induced 17-fold in galls relative to the rest of the root. Group II genes included *AtAAP1*, *AtAUX1*, *AtPIP2.6*, *At2g41190*, *AtMRP4*, and *AtLHT4* (Fig. 3). These genes were expressed at similar levels in galls and uninfected sections of root. Group III comprised genes that were expressed more highly in the surrounding root tissue compared with the galls. One gene was *AtANT1*, whose expression was significantly altered ($P = 0.004$). This gene was downregulated at all three timepoints according to the array results. Another gene, *At1g80510*, a close relative of *AtANT1*, was not significantly different between the two tissue types, but was upregulated at all three timepoints compared with the control in the array study.

Table 1. Overview of genes encoding transport proteins whose expression was significantly altered upon nematode infestation^a

Locus tag	Fold change(log ₂)			P value ^b	Pattern ^c	Name	Family ^d
	Week 1	Week 2	Week 4				
At1g58340	0.155	0.253	1.209	0.00099	UUU	AtDTX48	MOP
At2g21050	0.361	0.636	0.948	0.00109	UUU	AtAUX4/LAX3	AAAP
At5g13580	1.065	0.580	0.067	0.00138	UUU	AtWBC6	ABC
At5g57110	0.677	1.345	0.788	0.00146	UUU	AtACA8	P-ATPase
At4g39030	0.795	0.470	0.630	0.00179	UUU	AtDTX47	MOP
At5g38030	0.772	0.780	0.025	0.00256	UUU	AtDTX30	MOP
At2g39350	0.298	0.526	0.118	0.00305	UUU	AtWBC1	ABC
At2g38120	0.297	0.328	0.607	0.00339	UUU	AtAUX1	AAAP
At5g54800	0.600	0.457	0.068	0.00695	UUU	AtGPT1	DMT
At3g45680	1.652	1.014	0.223	0.00709	UUU	PTR	POT
At1g64780	1.399	0.565	0.350	0.00831	UUU	AtAMT1;2	AMT
At1g71880	0.965	0.823	0.853	0.00831	UUU	SUC1	GPH
At1g58360	0.960	0.922	0.042	0.00834	UUU	AtAAP1	AAAP
At2g39010	0.538	0.770	0.094	0.01110	UUU	AtPIP2.6	MIP
At5g44370	0.499	0.231	0.127	0.01211	UUU	Transporter	MFS
At3g51860	0.628	0.698	-0.164	0.00006	UUD	AtCAX3	CaCA
At5g12860	0.692	0.342	-0.080	0.00047	UUD	AtDiT1	DASS
At2g47800	0.501	0.689	-0.048	0.00633	UUD	AtMRP4	ABC
At1g76670	0.255	0.196	-0.041	0.00783	UUD	DMT	DMT
At1g72140	0.657	0.563	-0.003	0.00800	UUD	PTR	POT
At2g36910	0.472	0.481	-0.105	0.01013	UUD	AtMDR1	ABC
At5g27350	0.302	0.147	-0.124	0.01256	UUD	AtSFP1	MFS
At5g64290	0.304	0.146	-0.143	0.01278	UUD	AtDiT2.1	DASS
At3g03090	0.416	0.489	-0.078	0.01421	UUD	STP	MFS
At2g41560	0.497	0.217	-0.209	0.01606	UUD	AtACA4	P-ATPase
At1g20840	0.967	-0.082	0.497	0.01539	UDU	STP	MFS
At2g38290	0.116	-0.660	-0.622	0.00488	UDD	AtAMT2;1	AMT
At4g32650	0.094	-0.348	-1.036	0.00592	UDD	AtAKT4/AtKC1	VIC
At1g60960	0.067	-0.223	-1.027	0.00738	UDD	AtIRT3	ZIP
At3g12100	0.042	-0.213	-0.624	0.01209	UDD	AtMTPc2	CDF
At5g14940	-0.426	-0.422	-0.711	0.00003	DDD	PTR	POT
At1g69850	-0.415	-0.589	-0.485	0.00008	DDD	AtNRT1;2	POT
At3g43790	-0.325	-0.243	-0.439	0.00017	DDD	Transporter	MFS
At2g29650	-0.294	-0.471	-0.420	0.00037	DDD	Transporter	MFS
At3g17700	-0.145	-0.949	-0.873	0.00046	DDD	AtCNGC20	VIC
At3g60160	-0.534	-0.194	-0.482	0.00118	DDD	AtMRP9	ABC
At5g46040	-0.501	-0.670	-0.807	0.00140	DDD	PTR	POT
At5g13740	-0.362	-0.214	-0.480	0.00200	DDD	Transporter	MFS
At5g53550	-0.243	-0.515	-0.168	0.00282	DDD	AtYSL3	OPT
At2g40460	-0.308	-0.435	-0.626	0.00285	DDD	PTR	POT
At3g11900	-0.001	-0.441	-0.773	0.00411	DDD	AtANT1	AAAP
At2g30080	-0.215	-0.332	-0.654	0.00413	DDD	AtZIP6	ZIP
At4g31710	-0.213	-0.336	-0.621	0.00495	DDD	AtGLR2.4	GLR
At5g67330	-0.231	-0.006	-0.666	0.00606	DDD	AtNRAMP4	NRAMP
At5g52450	-0.466	-0.238	-0.517	0.00650	DDD	AtDTX16	MOP
At5g64840	-0.532	-0.455	-0.695	0.00759	DDD	ABC	ABC
At3g51870	-0.344	-0.374	-0.593	0.00832	DDD	MC	MC
At4g18210	-0.335	-0.660	-0.715	0.01252	DDD	AtPUP7	DMT
At1g59740	-0.153	-0.496	-0.783	0.01273	DDD	PTR	POT
At1g47530	-0.416	-0.536	-0.448	0.01531	DDD	AtDTX33	MOP

^a Transporter nomenclature is according to the ARAMEMNON database (Schwacke et al. 2003).

^b Analysis of variance.

^c Pattern refers to whether genes were upregulated (U) or downregulated (D) at week 1, 2, or 4 after infestation.

^d Family abbreviations taken from Saier (2000).

The tissue-specific expression pattern of a representative of the group I genes was demonstrated using transgenic plants expressing the *uidA* gene under control of the *AtAAP6* promoter 2 weeks after infestation (Fig. 4). The staining pattern of GUS activity in uninfested plants was consistent with a previous study (Okumoto et al. 2002), where it was localized to the vasculature of leaves and regions of lateral root branching (Fig. 4C and D, arrow heads). In infested plants, additional, stronger staining, which is consistent with the RT-PCR data, was observed in the galls of infested plants (Fig. 4C and D, arrows). To localize the expression of two group I genes (*AtAAP6* and *AtPIP2.5*) at higher resolution, thin sections were cut from the knots that had been imbedded in agarose. Analysis of these sections showed that *AtAAP6* and *AtPIP2.5* were expressed throughout the entire knot, including the giant cells (Fig. 4E and F).

DISCUSSION

Nematode-induced giant cells act as feeding sites and display transfer cell-like features (Offler et al. 2003). Previous studies identified two transporters (*AtSUC2* and *TobRB7*) as being involved in nematode feeding sites (Juergensen et al. 2003; Opperman et al. 1994). Recently, it was shown that *AtSUC2* is not expressed in syncytia (Hoth et al. 2005). To provide a global perspective of transporter gene expression during the infestation of *Arabidopsis* plants with the nematode *M. incognita*, a microarray analysis was completed. We found 50 transporter genes whose expression was significantly affected by nematode infestation. The transporter genes are representative of 19 of approximately 80 transporter gene families in *Arabidopsis* (Busch and Saier 2002). For the majority of the transporters that were regulated in this study (38/50), this is the first report providing any evidence of a connection to some biological process, such as involvement in nematode feeding sites or response to nematode infestation. The representation of specific families of genes in the list of 50 varied between 6 and 17%. For example, the *AtCAX* family has six genes on the GeneChip and the expression of one was significantly altered, which represents approximately 17% of the genes in that family. For two large gene families which we quantified, the percentage of genes expressed in specific family members present on the GeneChip, such as ATP binding cassette transporters

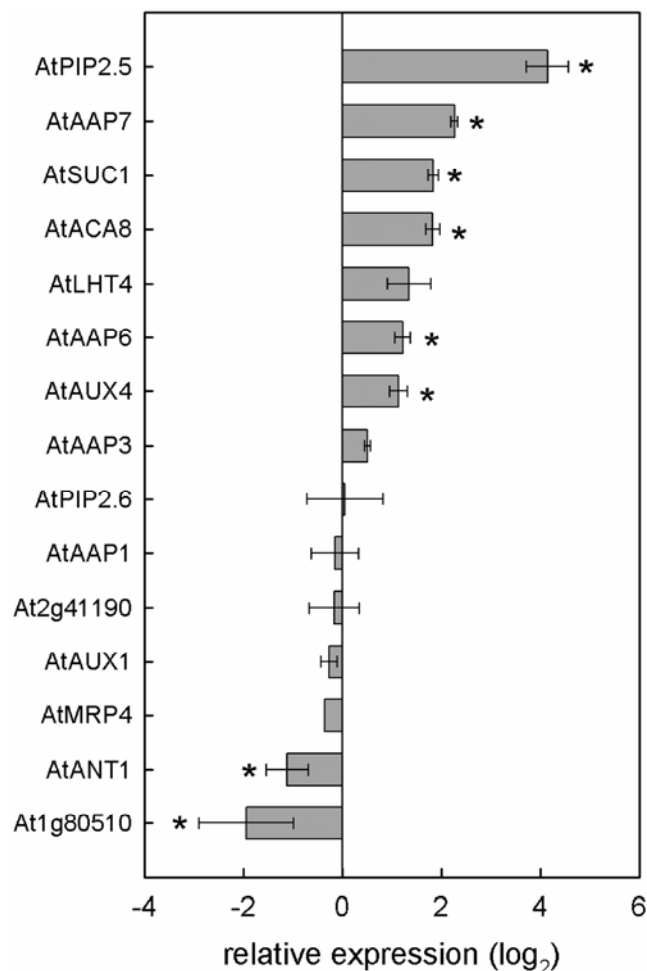


Fig. 3. Expression of 15 genes in knots compared with the rest of the infested roots quantified using real-time reverse-transcriptase polymerase chain reaction. Knots were excised manually from infested roots 2 weeks after infestation. The relative expression levels (log₂) were calculated in three reps by dividing expression levels in knots with expression levels in the uninfested regions of the roots. Positive values indicate higher expression in the knot and negative values indicate higher expression in the rest of the root. Asterisks indicate a significant difference between expression levels in the knot compared with the rest of the root ($P < 0.05$).

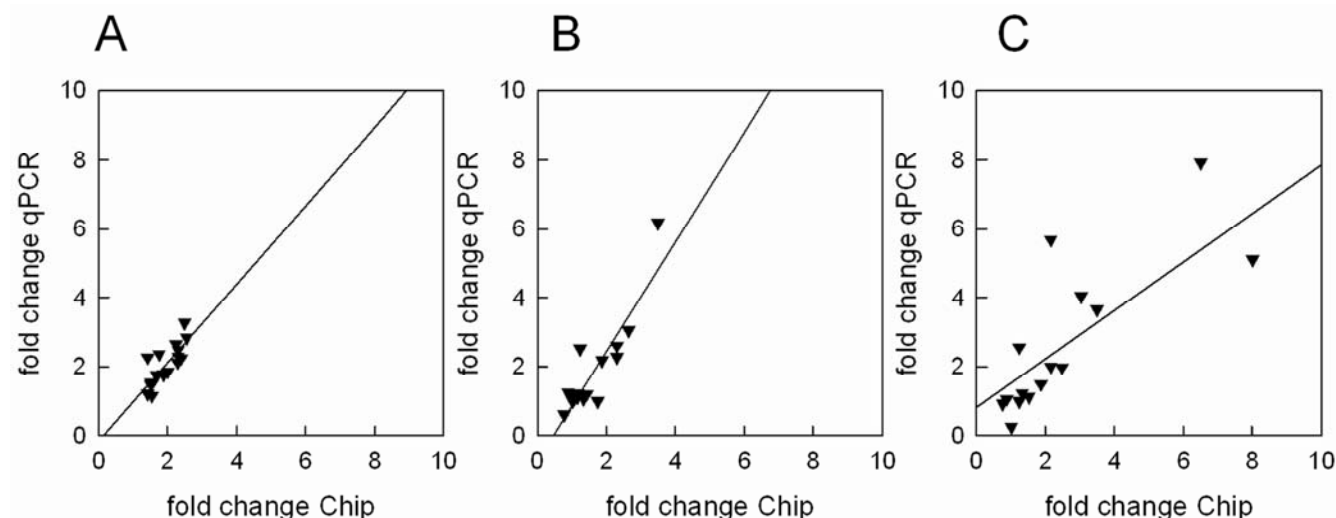


Fig. 2. Correlation between fold changes measured with GeneChip and quantitative real-time reverse-transcriptase polymerase chain reaction (RT-PCR) of 16 genes. The same RNA used to hybridize the arrays was used for real time RT-PCR to determine the reproducibility of the fold changes. Significant correlations between the two methods were obtained in all timepoints. The linear regression line is shown. **A**, Week 1 ($r^2 = 0.62$, $P = 0.0004$); **B**, week 2 ($r^2 = 0.78$, $P < 0.0001$); **C**, week 4 ($r^2 = 0.62$, $P < 0.0001$).

(ABC) (102 family members on GeneChip) and the AAAP family (46 family members on GeneChip), representation was relatively low (6 and 9%, respectively). In the case of the peptide transporter family genes (52 family members on GeneChip), representation was relatively high (13%).

Transporters in symbiosis and pathogenesis.

Transport processes are known to be important in interactions between plants and other living organisms. In two of the most widespread symbiotic interactions with fungi and bacteria, transporters for phosphate (Harrison et al. 2002; Rausch et al. 2001) and sugars (Flemetakis et al. 2003), channels with conductivities for a number of ions, including ammonium (Tyerman et al. 1995) and aquaporins (Krajinski et al. 2000), have been suggested to play an important role in transfer of minerals and carbon between plants and symbiotic partners. A

putative channel-like protein recently was implicated in the ability of nitrogen-fixing bacteria to infect the legume *M. truncatula* (Ane et al. 2004).

In contrast to what is known about the functioning of membrane transport proteins in symbiotic interactions, less is known about the role membrane transport proteins play in plant–parasite interactions. In one case, the disruption of a cyclic nucleotide gated channel has been shown to confer broad-spectrum disease resistance to a bacterial pathogen (Clough et al. 2000). Genes encoding sugar and water transporters have been found to be induced upon wounding and fungal pathogen interaction (Fotopoulos et al. 2003; Meyer et al. 2004; Truernit et al. 1996; Werner et al. 2001). Upon infestation by the nematode *M. incognita*, we found that the expression of 50 transport proteins was significantly different between the nematode-infested roots and the control. This focused analysis of GeneChip data

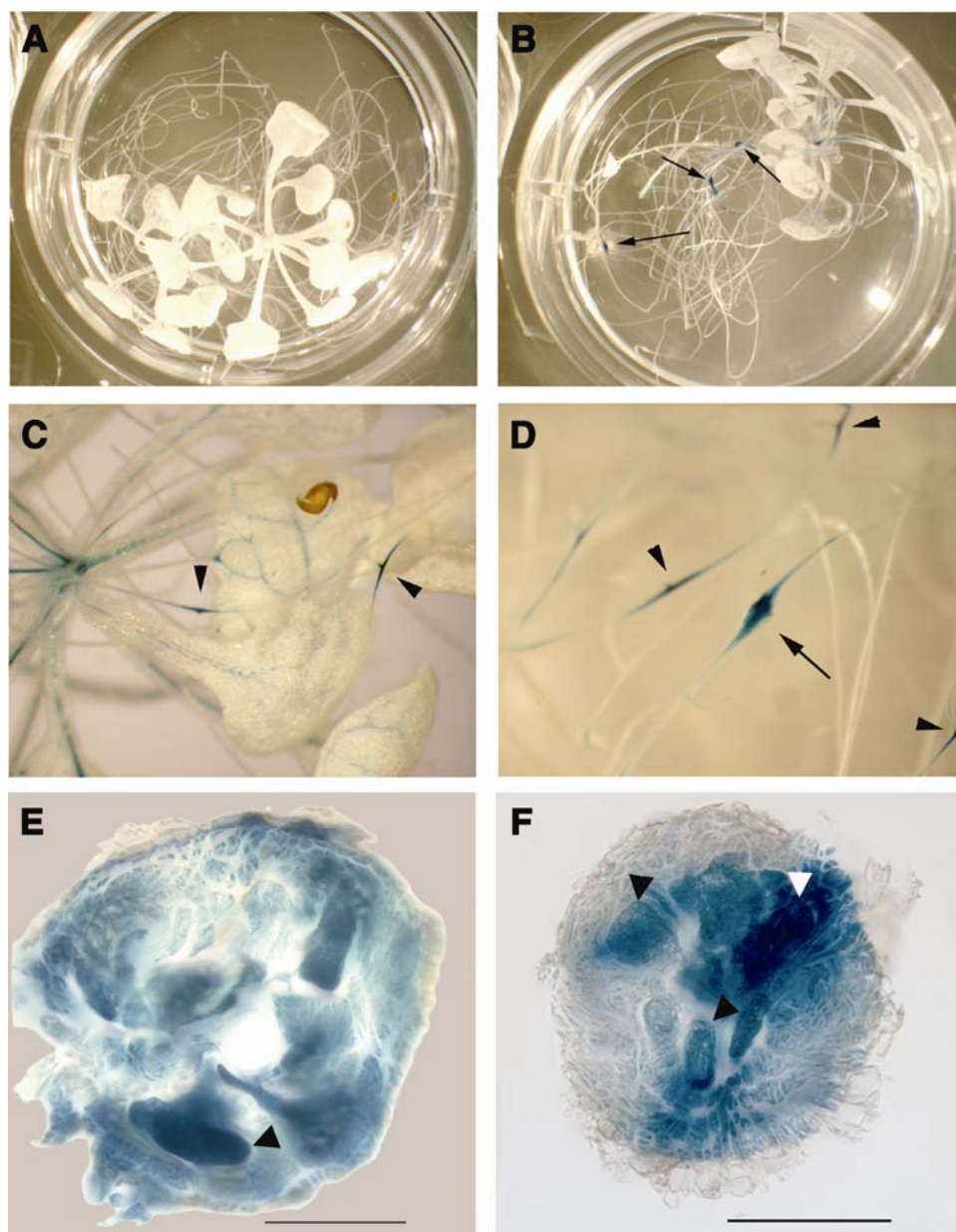


Fig. 4. Expression of the *uidA* gene under control of the *AtAAP6* or *AtPIP2.5* promoter. Gene expression in **B**, **D**, **E**, and **F**, plants 2 weeks after infestation and **A** and **B**, uninfested control plants. The expression level of *AtAAP6* is higher in knots (**B**, arrows) than in the uninfested roots (**A**). In addition to the knots (**D**, arrow), the gene is expressed in the region of root branching (**C** and **D**, arrowheads) in both uninfested and infested roots. Thin sections of nematode-infested knots **E**, *AtAAP6* promoter::GUS and **F**, *AtPIP2.5* promoter::GUS. Staining is visible throughout most of the knot, including the giant cells. Arrowheads indicate the location of some of the giant cells. Bars = 100 μ M.

provides a comprehensive analysis of transporter gene expression in a plant–parasite interaction. We also analyzed the supplemental data from a previous study using the Affymetrix 8K GeneChip which focused on the early response to infestation by the beet cyst nematode *Heterodera schachtii*. From those data, we found that only seven transporter genes were induced (Puthoff et al. 2003). Two of these, *AtAAP6* and *AtPIP2.5*, also were found to be induced by *M. incognita* and, therefore, may be involved in similar processes in the different types of feeding sites. The remaining 48 genes encoding transporters found in our study were not identified in any previous study on nematodes.

Giant-cell development.

Data from our electron microscopy study in *Arabidopsis* support findings from root-knot nematode interactions in other plant species (Jones and Payne 1978). In *Arabidopsis*, we observed many features common to giant cells found in other plant species. Interestingly, two dynamic aspects of giant-cell formation were the observed changes in vacuolar components of the developing giant cell and the development of cell wall ingrowths. When giant cells are initiated, they contain vacuolar structures with size and distribution similar to adjacent nongiant cells. Giant cells at intermediate stages of development prominently display an expanded central vacuole while they maintain smaller vacuolar structures in the peripheral regions of the cell. At present, it is not clear whether these two populations of vacuoles are functionally identical (Bassham and Raikhel 2000; Bethke and Jones 2000). Therefore, the derivation of the large vacuole found in giant cells of intermediate development is currently unknown (Fig. 1B). Still unexplained are the processes that govern the disappearance of this large vacuole as the giant cell matures (Fig. 1C).

Our studies revealed large changes in the ultrastructure of the giant cells over time in *Arabidopsis* roots, resulting in giant cells with a wall labyrinth, which is the hallmark of a transfer cell (Offler et al. 2003). Transfer cells occur in a variety of tissues at bottlenecks within nutrient transport pathways (Offler et al. 2003). Cell wall proliferation is evident at intermediate stages in giant-cell development (Fig. 1C). However, formation of a cell wall labyrinth occurs only at late stages in development. As the giant cells developed, we also observed many changes in transporter gene expression. In some cases, transporters may play a role in giant-cell development or may transport nutrients into giant cells that are required for nematodes.

Transporters found in developing feeding sites.

Previous studies of transport processes in transfer-like cells focused on specific transport proteins such as H^+ -ATPases, sucrose, and amino acid transporters (Offler et al. 2003). The use of the ATH1 GeneChip provided a comprehensive picture of the transport processes induced in *Arabidopsis* roots during the development of nematode feeding sites. The three timepoints chosen for the GeneChip analysis represent three phases of giant-cell development: initiation, establishment and swelling of giant cells, and maintenance. Because it was not possible to synchronize nematode infestation at the start of these experiments, samples represent the developmental stage which the majority of cells in the knots had reached at each timepoint.

The earliest timepoint which corresponds to the induction of feeding sites was characterized by the expression of organelle-specific transporters and primary active transport systems. This may indicate an elevated energy demand by the root, which is consistent with earlier studies (Gheysen and Fenoll 2002; McClure 1977). Two members of the Ca^{2+} -ATPase (ACA) family were upregulated in nematode-infested roots. *AtACA4* has

been localized to small vacuolar structures which are abundant at early stages of development (Geisler et al. 2000). *AtACA8* has been localized to the plasma membrane (Bonza et al. 2000). The expression of *AtACA8* was higher in feeding sites than in the rest of the root, which may be indicative of a more specific role in giant-cell formation, but additional data are needed to test this hypothesis. These two genes and *AtCAX3* (Hirschi 2001), a member of the Ca^{2+} :cation antiporter family, have been suggested to play a role in modulating Ca^{2+} -mediated signaling events (Sanders et al. 1999). Based on these data, we suggest that Ca^{2+} influx and signaling play a role in plant response to nematode infestation and possibly giant-cell or transfer-cell induction and establishment.

At the intermediate timepoint, 2 weeks after infestation, some of the key features of giant-cell development were rapid cell expansion, formation of a large central vacuole, and increased deposition of cell wall material. Changes in expression patterns of some membrane transport proteins in nematode-infested roots could reflect changes in gene expression in giant cells or in uninfested regions of the root. Therefore, we compared mRNA isolated from dissected root-knots at week 2 with regions of the root that were not infested. We found one aquaporin gene, *AtPIP2.6*, to be significantly induced in infested roots. According to its expression pattern, this gene appears to be induced throughout infested roots with no significant tissue specificity. In contrast, the expression of its closest relative, *AtPIP2.5*, which also is induced upon infestation with beet cyst nematode *H. schachtii* (Puthoff et al. 2003), was 17-fold higher in root-knot tissue compared with the rest of the root. The localization of *AtPIP2.5* to the root-knot tissue suggests that this gene may be a functional ortholog of a giant-cell-specific aquaporin TobRB7 from tobacco (Opperman et al. 1994; Puthoff et al. 2003). More detailed localization showed that this aquaporin is expressed throughout the gall, including in giant cells.

A sucrose transporter gene, *AtSUC1*, was significantly induced upon root-knot nematode infestation. The expression of this gene also was higher in feeding sites than in the surrounding tissue (Fig. 3), which is consistent with what is known about the expression of sucrose transporters in seed coat transfer cells (Harrington et al. 1997; Wang et al. 1995). In giant cells, sucrose may be a major osmolyte driving swelling and *AtSUC1* may be partly responsible for sucrose transport into the developing giant cells. In contrast, the sucrose transporter gene *AtSUC2* was found to be upregulated in syncytia (Juergensen et al. 2003), but does not appear to be directly involved in the active loading of sucrose into syncytia (Hoth et al. 2005). The expression of this transporter gene was unaltered in our experiments. The expression of different sucrose transporters in cyst and root-knot nematode-infested *Arabidopsis* roots provides additional support for the conclusion that syncytia and giant-cell formation follow different developmental programs.

Amino acid transport in feeding sites.

We observed large changes in the expression of genes involved in amino acid transport. Overall, the changes were a downregulation of peptide transport and an upregulation of amino acid transporters. In contrast, more peptide transporters than amino acid transporters were induced by cyst nematodes (Puthoff et al. 2003). Most of the amino acid transporters that we investigated in detail were expressed at similar levels in knots versus the rest of the root, suggesting a role for these transporters in the increased amino acid transport capacity of infested roots (Fischer et al. 1995; Okumoto et al. 2002). The expression of *AtAAP6* was characterized by both real-time PCR and a *promoter::GUS* assay. Real-time RT-PCR analysis demonstrated that *AtAAP6* was expressed to higher levels in

knots. This finding was supported by an *AtAAP6 promoter::GUS* study which showed that the activity of this promoter was much higher in galls compared with the surrounding tissue and was expressed in both giant cells and other cells in the galls. The role that the amino acid transporter *AtAAP6* plays in both cysts and root knots is not yet defined.

Two genes that have been grouped into the AAP superfamily were induced at the late timepoint. Those genes were *AtAUX1* and *AtAUX4/LAX3* (At 2g2150), which encode putative auxin transporters (Bennett et al. 1996; Parry et al. 2001; Swarup et al. 2001). Auxin is required for formation of syncytia (Goverse et al. 2000), and auxin levels increase in plant roots infested with nematodes (Viglierchio and Yu 1968). Further studies are required to confirm the substrates that the AUX proteins transport and to determine their role in the establishment and maintenance of the nematode feeding sites.

The results from the microarray experiments provide a global overview of the regulation of 634 transporter genes in the *Arabidopsis*–root-knot nematode interaction. Further work will be needed to determine the role and functional characteristics of the many newly identified genes that encode transporters regulated by nematode infestation.

MATERIALS AND METHODS

Sample preparation for microscopy.

M. incognita-infested roots of *Arabidopsis thaliana* (Col-0), grown as described below, were the source of tissues for electron microscopy. Root sections containing giant cells were dissected to fit the specimen planchettes for high-pressure freezing; they were packed in an aqueous solution of 150 mM mannitol and frozen in a Bal-Tec high-pressure freezer. Samples were freeze substituted in acetone containing 2% osmium tetroxide for 3 days at -85°C , 15 h at -60°C , and 5 h at -30°C . Then, they were warmed gradually to room temperature, rinsed in acetone, and slowly, over a period of 7 days, infiltrated with Epon-Araldite resin. Thin sections were stained in uranyl and lead salts and observed in a Leo 912 transmission electron microscope (TEM); images were collected with a digital camera.

Microarray sample preparation.

Wild-type *Arabidopsis* Col-0 seed were sterilized and germinated on agar plates (2% sucrose, 0.3% Gamborg's B-5, 0.6% Gelrite, pH 6.1, 8 h of light and 16 h of darkness, 23°C). After 7 days, five seedlings were transferred onto each of 84 plates and grown vertically for 3 weeks. At 3 weeks after germination, approximately 1,000 root-knot nematode (*M. incognita*) eggs were added to half of the plates. A set of uninfested plates was kept as a control group. Roots from 70 plants (14 plates containing 5 plants each) were pooled and composed one replicate sample. At each timepoint, three replicates were harvested from both infested and noninfested plants at 1, 2, and 4 weeks after inoculation. Infested plants had at least 10 galls per root. RNA was extracted using the RNeasy mini-kit protocol (Qiagen, Valencia, CA, U.S.A.) following the manufacturer's instructions. Isolated RNA was converted into cRNA as described by the Affymetrix protocol. Briefly, cDNAs were synthesized from 20 μg of the extracted RNA with the SuperScript Choice System (Invitrogen, Carlsbad, U.S.A.). The BioArray HighYield RNA transcript labeling kit (Enzo Biochem, New York) then was used to transcribe and label the cRNA. The cRNA was fragmented to 200 nucleotides or less, heated at 99°C for 5 min, and hybridized for 16 h at 45°C to *Arabidopsis* ATH1 Genome Arrays (Affymetrix Inc., Santa Clara, CA, U.S.A.). The fluorescence was visualized with an Agilent confocal laser scanner.

Microarray analysis.

Image data was quantified using GeneChip Analysis Suite/Microarray Suite 5.0 (MAS 5.0). Quality control checks were performed according to standard Affymetrix protocols. In addition, we assessed the concordance among replicates using a weighted kappa coefficient (Fleiss 1981). All replicates were found to be of high quality and so we proceeded with the analyses.

The identification of informative features was performed using default settings (α_1 : 0.04; α_2 : 0.06; δ : 0.015; scale factor: 1.0; norm factor: 1.0). If all replicates for a particular feature were deemed "absent," that feature was removed from further consideration. We used the AMPL database (Ward 2001) to identify known transporters. There were 850 transporters identified in this database at the time of our search. We used this list to subset the ATH1 chip and we only analyzed features which corresponded to the known transporters. The features identified as transporters ($n = 634$) were analyzed. Transcript levels were normalized to the overall chip median and log transformed. For each feature, which represents the combined expression data from all relevant probe pairs on the chip, the effect of the mutant and the time stimulus were examined in an analysis of variance modeling framework. (Kerr and Churchill 2001a and b; Oleksiak et al. 2002; Singh et al. 2003; Wayne and McIntyre 2002; Wolfinger et al. 2001). The model $Y_{ijkl} = \mu + \psi_i + \omega_j + \psi\omega_{ij} + \epsilon_{ijkl}$ was fit, where Y is the normalized and log transformed estimate of signal intensity. The parameter μ was the overall mean intensity for that feature, ψ represents the effect of the condition (nematode versus control), and ω represents the effect of week ($j = 1, 2, \text{ or } 4$). To test the null hypothesis that a particular feature's expression level was not different across conditions, an F test of the effect of condition was conducted for each feature and a P value was calculated. We examined the model for conformation to the assumption of normality of the residuals by testing the null hypothesis so that the residuals for each feature were normally distributed using the Shapiro-Wilkes test. To determine whether a feature was statistically significant for the effect of condition, we used a false discovery rate (FDR) (Benjamini and Hochberg 1995) of 0.20 for each of the two analyses. Type I and type II errors are inversely related, with decreases in false positives (type I) being associated with increases in false negatives (type II); therefore, we chose an FDR of 0.2 in order to minimize the type II errors without disregarding the type I error. When many tests are performed, keeping the proportion of false discoveries relative to all significant results at a low level is a powerful alternative to the traditional approach of avoiding even a single false discovery (family-wise error rate [FWER]). Control of the FWER at α , via Bonferroni or sequential Bonferroni procedures (Holm 1979), is a suitable approach only if the penalty of making even one type I error is severe. In many studies, including this one, avoiding any type I error irrespective of its cost in terms of type II errors is not a satisfactory approach. The Benjamini and Hochberg procedure for FDR control provides a sensible solution: it offers an easily interpretable mechanism to control type I errors while simultaneously reducing type II errors. Control of the false discovery rate is becoming widely used in genomic research (Peng et al. 2003; Sabatti et al. 2003; Storey and Tibshirani 2003; Tusher et al. 2001; Veena et al. 2003; Verhoeven et al. 2005; Weller et al. 1998). Microarray analyses were performed using SAS software (SAS Institute, Cary, NC, U.S.A.).

Data access.

The processed microarray data are presented according to MIAME standards (Brazma et al. 2001).

RNA isolation and cDNA synthesis for localization studies.

Roots were harvested from plants 2 weeks postinfestation. The week 2 timepoint was chosen for this analysis because this is the first time period at which knots are readily excised from the surrounding tissue. Knots were excised with a scalpel and the remaining tissue was separated. The two samples were frozen immediately in liquid nitrogen and RNA from both tissues was isolated independently using the QIAGEN RNA isolation kit (Qiagen) following the manufacturer's instructions.

RNA was treated with DNaseI (AMBION, Austin, TX, U.S.A.) according to the manufacturer's protocol. cDNA was synthesized using the Eppendorf cMaster cDNA Synthesis Kit (Eppendorf, Westbury, NY, U.S.A.) and Oligo-dT priming following the manufacturer's protocol.

Quantitative PCR.

Primers flanking a 90- to 320-bp region specific to the last 800 bases of the 3' region of each gene of interest and two internal standards (UBQ 10 At4g05320 and eIF4 At3g60240) were designed using PrimerSelect (DNASTar, Lasergene, Madison, WI, U.S.A.). Real-time PCR was performed by using an iCycler (BioRAD, Hercules, CA, U.S.A.) and SyBr green I (Molecular Probes, Eugene, OR, U.S.A.) detection. All reactions were carried out in triplicates. The reactions contained 20 nM Fluorescein (BioRAD) as a passive reference dye. PCR was carried out using the heat-activated Hot Master Taq polymerase (Eppendorf) spiked with 1 µl of a 1:10,000 dilution of SYBR green I. The amount of targeted messages in each sample was determined by relating the iCycler results for the gene of interest to a standard curve for each individual gene. Expression levels were normalized to both UBQ10 and eIF4 to ensure the similarity of relative expression levels.

For the correlation between chip and RT-PCR results, a linear regression model $y = y_0 + ax$ was fit and examined for goodness of fit using the r^2 value.

To test whether a gene was expressed in a specific tissue at a higher level, the null hypothesis that a particular gene's expression level was not different in the two tissues, a t test of the effect of condition was conducted for each gene and a P value was calculated. When $P < 0.05$, the null hypothesis was rejected.

Cloning of the promoter fragments and staining for GUS activity.

The *AtAAP6 promoter::uidA* fusion construct was described previously (Okumoto et al. 2002). The *AtPIP2.5* promoter fragment was amplified from Col-0 genomic DNA using the oligonucleotides 5'-GTCTTCTTATTCTTCACCAAACCTTAAG-3' and 5'-GAGGAATAAATTAACAGATTATCACAC-3'. The resulting product was cloned in front of the *uidA* gene as a transcriptional fusion. For histochemical localization of GUS activity, plant material was infiltrated with X-Gluc at 1 mg/ml in staining buffer (100 mM Na-Phosphate, pH 7.0, 10 mM EDTA, 10 mM $K_4[Fe_2(CN)_6]$, 10 mM $K_3[Fe_2(CN)_6]$, 1% Triton X-100, and 10% dimethyl sulfoxide). Staining was performed for 7 h at 37°C. After staining was completed, the plants were transferred to 70% (vol/vol) ethanol. For thin sections, root material was fixed overnight in 3% (vol/vol) glutaraldehyde containing 100 mM Tris, pH 7.5, and 0.03% Triton-X 100. Roots then were imbedded in 3% low-melting-point agarose. Specimens were sectioned to 75 µm using a Vibratome Series 1500.

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AUTHOR-RECOMMENDED INTERNET RESOURCES

Arabidopsis Membrane Protein Library (AMPL) database
www.cbs.umn.edu/arabidopsis/index2.htm
 Aramemnon plant membrane protein database:
aramemnon.botanik.uni-koeln.de
 Affymetrix protocol www.affymetrix.com/support/index.html.
 MIAME standards: www.mged.org/Workgroups/MIAME/miame.html
 TCDB Transport classification database: www.tcdb.org